desmosterolosis

Desmosterolosis is a condition that is characterized by neurological problems, such as brain abnormalities and developmental delay, and can also include other signs and symptoms.

Children with desmosterolosis have delayed speech and motor skills (such as sitting and walking). Later in childhood, some affected individuals are able to walk with support; verbal communication is often limited to a few words or phrases. Common brain abnormalities in desmosterolosis include malformation of the tissue that connects the left and right halves of the brain (the corpus callosum) and loss of white matter, which consists of nerve fibers covered by a fatty substance called myelin.

People with desmosterolosis commonly have muscle stiffness (spasticity) and stiff, rigid joints (arthrogryposis) affecting their hands and feet. Other features seen in some affected individuals include short stature, abnormal head size (either larger or smaller than normal), a small lower jaw (micrognathia), an opening in the roof of the mouth (cleft palate), involuntary eye movements (nystagmus) or eyes that do not look in the same direction (strabismus), heart defects, and seizures.

Frequency

The prevalence of desmosterolosis is unknown; at least 10 affected individuals have been described in the scientific literature.

Genetic Changes

Desmosterolosis is caused by mutations in the *DHCR24* gene. This gene provides instructions for making an enzyme called 24-dehydrocholesterol reductase, which is involved in the production (synthesis) of cholesterol. Cholesterol is a waxy, fat-like substance that can be obtained from foods that come from animals (particularly egg yolks, meat, poultry, fish, and dairy products). It can also be produced in various tissues in the body. For example, the brain cannot access the cholesterol that comes from food, so brain cells must produce their own. Cholesterol is necessary for normal embryonic development and has important functions both before and after birth.

DHCR24 gene mutations lead to the production of 24-dehydrocholesterol reductase with reduced activity. As a result, there is a decrease in cholesterol production. Because the brain relies solely on cellular production for cholesterol, it is most severely affected. Without adequate cholesterol, cell membranes are not formed properly and nerve cells are not protected by myelin, leading to the death of these cells. In addition, a decrease in cholesterol production has more severe effects before birth than during other periods of development because of the rapid increase in cell number that takes

place. Disruption of normal cell formation before birth likely accounts for the additional developmental abnormalities of desmosterolosis.

Inheritance Pattern

This condition is inherited in an autosomal recessive pattern, which means both copies of the gene in each cell have mutations. The parents of an individual with an autosomal recessive condition each carry one copy of the mutated gene, but they typically do not show signs and symptoms of the condition.

Other Names for This Condition

deficiency of 3beta-hydroxysterol delta24-reductase

Diagnosis & Management

Genetic Testing

 Genetic Testing Registry: Desmosterolosis https://www.ncbi.nlm.nih.gov/qtr/conditions/C1865596/

General Information from MedlinePlus

- Diagnostic Tests https://medlineplus.gov/diagnostictests.html
- Drug Therapy https://medlineplus.gov/drugtherapy.html
- Genetic Counseling https://medlineplus.gov/geneticcounseling.html
- Palliative Care https://medlineplus.gov/palliativecare.html
- Surgery and Rehabilitation https://medlineplus.gov/surgeryandrehabilitation.html

Additional Information & Resources

MedlinePlus

- Health Topic: Cholesterol https://medlineplus.gov/cholesterol.html
- Health Topic: Lipid Metabolism Disorders https://medlineplus.gov/lipidmetabolismdisorders.html

Genetic and Rare Diseases Information Center

 Desmosterolosis https://rarediseases.info.nih.gov/diseases/10283/desmosterolosis

Educational Resources

- American Heart Association: About Cholesterol http://www.heart.org/HEARTORG/Conditions/Cholesterol/AboutCholesterol/About-Cholesterol_UCM_001220_Article.jsp
- Boston Children's Hospital: Arthrogryposis http://www.childrenshospital.org/conditions-and-treatments/conditions/ arthrogryposis
- Centers for Disease Control and Prevention: Facts About Developmental Disabilities
 https://www.cdc.gov/ncbddd/developmentaldisabilities/facts.html
- Cleveland Clinic: Cholesterol http://my.clevelandclinic.org/health/articles/cholesterol
- Cleveland Clinic: Spasticity http://my.clevelandclinic.org/health/articles/spasticity
- Disease InfoSearch: Desmosterolosis http://www.diseaseinfosearch.org/Desmosterolosis/2221
- Kennedy Krieger Institute: Neurological Disorders
 https://www.kennedykrieger.org/patient-care/diagnoses-disorders/neurological-disorders
- KidsHealth from Nemours: Cholesterol and Your Child http://kidshealth.org/en/parents/cholesterol.html
- KidsHealth from Nemours: Delayed Speech or Language Development http://kidshealth.org/en/parents/not-talk.html
- MalaCards: desmosterolosis http://www.malacards.org/card/desmosterolosis
- Merck Manual Home Edition for Patients and Caregivers: Overview of Cholesterol and Lipid Disorders
 http://www.merckmanuals.com/home/hormonal-and-metabolic-disorders/ cholesterol-disorders/overview-of-cholesterol-and-lipid-disorders
- Orphanet: Desmosterolosis http://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=35107

Patient Support and Advocacy Resources

- American Association on Intellectual and Developmental Disabilities (AAIDD) http://aaidd.org/
- National Organization for Disorders of the Corpus Callosum http://nodcc.org/

- Resource List from the University of Kansas Medical Center: Developmental Delay/ Mental Retardation
 - http://www.kumc.edu/gec/support/devdelay.html
- Sterol & Isoprenoid Research Consortium http://www.rarediseasesnetwork.org/cms/STAIR

Scientific Articles on PubMed

PubMed

https://www.ncbi.nlm.nih.gov/pubmed?term=%28desmosterolosis%5BTIAB%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D

OMIM

 DESMOSTEROLOSIS http://omim.org/entry/602398

Sources for This Summary

- Dias C, Rupps R, Millar B, Choi K, Marra M, Demos M, Kratz LE, Boerkoel CF. Desmosterolosis: an illustration of diagnostic ambiguity of cholesterol synthesis disorders. Orphanet J Rare Dis. 2014 Jun 25;9:94. doi: 10.1186/1750-1172-9-94.
 - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/24961299
 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4076431/
- Kanungo S, Soares N, He M, Steiner RD. Sterol metabolism disorders and neurodevelopment-an update. Dev Disabil Res Rev. 2013;17(3):197-210. doi: 10.1002/ddrr.1114. Review.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/23798009
- Schaaf CP, Koster J, Katsonis P, Kratz L, Shchelochkov OA, Scaglia F, Kelley RI, Lichtarge O, Waterham HR, Shinawi M. Desmosterolosis-phenotypic and molecular characterization of a third case and review of the literature. Am J Med Genet A. 2011 Jul;155A(7):1597-604. doi: 10.1002/ajmg.a.34040. Epub 2011 Jun 10. Review.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/21671375
- Waterham HR, Koster J, Romeijn GJ, Hennekam RC, Vreken P, Andersson HC, FitzPatrick DR, Kelley RI, Wanders RJ. Mutations in the 3beta-hydroxysterol Delta24-reductase gene cause desmosterolosis, an autosomal recessive disorder of cholesterol biosynthesis. Am J Hum Genet. 2001 Oct;69(4):685-94. Epub 2001 Aug 22.
 Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/11519011
 Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1226055/
- Zolotushko J, Flusser H, Markus B, Shelef I, Langer Y, Heverin M, Björkhem I, Sivan S, Birk OS.
 The desmosterolosis phenotype: spasticity, microcephaly and micrognathia with agenesis of
 corpus callosum and loss of white matter. Eur J Hum Genet. 2011 Sep;19(9):942-6. doi: 10.1038/ejhg.2011.74. Epub 2011 May 11.

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